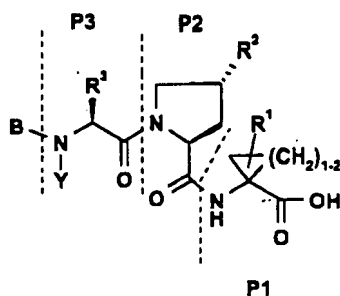


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wherein **B** is H, a C₆ or C₁₀ aryl, C₇₋₁₈ aralkyl; Het or (lower alkyl)-Het, all of which optionally substituted with C₁₋₆ alkyl; C₁₋₈ alkoxy; C₁₋₈ alkanoyl; hydroxy; hydroxyalkyl; halo; haloalkyl; nitro; cyano; cyanoalkyl; amino optionally substituted with C₁₋₈ alkyl; amido; or (lower alkyl)amide;

or **B** is an acyl derivative of formula **R₄-C(O)-**; a carboxyl derivative of formula **R₄-O-C(O)-**; an amide derivative of formula **R₄-N(R₅)-C(O)-**; a thioamide derivative of formula **R₄-N(R₅)-C(S)-**; or a sulfonyl derivative of formula **R₄-SO₂** wherein

R₄ is (i) C₁₋₁₀ alkyl optionally substituted with carboxyl, C₁₋₈ alkanoyl, hydroxy, C₁₋₈ alkoxy, amino optionally mono- or di-substituted with C₁₋₈ alkyl, amido, or (lower alkyl) amide;

(ii) C₃₋₇ cycloalkyl, C₃₋₇ cycloalkoxy, or C₄₋₁₀ alkylcycloalkyl, all optionally substituted with hydroxy, carboxyl, (C₁₋₆ alkoxy)carbonyl, amino optionally mono- or di-substituted with C₁₋₈ alkyl, amido, or (lower alkyl) amide;

(iii) amino optionally mono- or di-substituted with C₁₋₈ alkyl; amido; or (lower alkyl)amide;

(iv) C₆ or C₁₀ aryl or C₇₋₁₈ aralkyl, all optionally substituted with C₁₋₆ alkyl, hydroxy, amido, (lower alkyl)amide, or amino optionally mono- or di-substituted with C₁₋₈ alkyl; or

(v) Het or (lower alkyl)-Het, both optionally substituted with C₁₋₆ alkyl, hydroxy, amido, (lower alkyl) amide, or amino optionally mono- or di-substituted with C₁₋₈ alkyl;

R₅ is H or C₁₋₈ alkyl; with the proviso that when **B** is a carboxyl derivative, an amide derivative or a thioamide derivative, **R₄** is not a cycloalkoxy;

Y is H or C₁₋₆ alkyl;

R² is C₁₋₈ alkyl, C₃₋₇ cycloalkyl, or C₄₋₁₀ alkylcycloalkyl, all optionally substituted with hydroxy, C₁₋₈ alkoxy, C₁₋₈ thioalkyl, amido, (lower alkyl)amido, C₆ or C₁₀ aryl, or C₇₋₁₈

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aralkyl;

R² is CH₂-R₂₀, NH-R₂₀, O-R₂₀ or S-R₂₀, wherein R₂₀ is pyridinyl, quinolyl, (lower alkyl)-pyridinyl or (lower alkyl)-quinolyl, each optionally mono-, di- or tri-substituted with R₂₁,

wherein each R₂₁ is independently C₁₋₆ alkyl; C₁₋₆ alkoxy; lower thioalkyl; sulfonyl;

NO₂; OH; SH; halo; haloalkyl; amino optionally mono- or di-substituted with C₁₋₆

alkyl, C₆ or C₁₀ aryl, C₇₋₁₄ aralkyl, Het or (lower alkyl)-Het; amido optionally mono-

substituted with C₁₋₆ alkyl, C₆ or C₁₀ aryl, C₇₋₁₄ aralkyl, Het or (lower alkyl)-Het;

carboxyl; carboxy(lower alkyl); C₆ or C₁₀ aryl, C₇₋₁₄ aralkyl or Het, said aryl, aralkyl

or Het being optionally substituted with R₂₂;

wherein R₂₂ is C₁₋₆ alkyl; C₃₋₇ cycloalkyl; C₁₋₆ alkoxy; amino optionally

mono- or di-substituted with C₁₋₆ alkyl; sulfonyl; (lower alkyl)sulfonyl; NO₂;

OH; SH; halo; haloalkyl; carboxyl; amide; (lower alkyl)amide; or Het

optionally substituted with C₁₋₆ alkyl;

R³ is H; C₁₋₆ alkyl, C₃₋₇ cycloalkyl, C₂₋₆ alkenyl, or C₂₋₆ alkynyl, all optionally substituted with halogen;

or a pharmaceutically acceptable salt or ester thereof;

wherein "Het" is defined as a five-membered saturated or unsaturated, aromatic or non-aromatic, heterocycle containing from one to four heteroatoms selected from nitrogen, oxygen and sulfur, wherein said heterocycle is optionally fused to a benzene ring.